

GGDEF domain as spatial on-switch for a phosphodiesterase by interaction with landmark protein HubP

- SUPPLEMENTARY INFORMATION -

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Supplementary Table 1: Bacterial strains that were used in this study

Strain	Genotype	Purpose	Reference
<i>Escherichia coli</i>			
DH5α λpir	φ80d ^r /lacZ ΔM15 Δ(lacZYA-argF)U169 recA ₁ hsdR17 deoR thi-1 supE44 gyrA96 relA/λpir	cloning strain	1
WM3064	thrB1004 pro thi rpsL hsdS lacZ ΔM15 RP4-1360 Δ(araBAD) 567ΔdapA 1341::[erm pir(wt)]	conjugation strain for <i>Shewanella</i>	2
BL21(DE3)	fhuA2 [lon] ompT gal (λ DE3) [dcm] ΔhsdS λ DE3 = λ sBamH1o ΔEcoRI-B int:(lacI::PlacUV5::T7 gene1) i21 Δnin5	protein overproduction strain	3
<i>Shewanella putrefaciens</i> CN-32			
S757	CN-32 wt	wildtype strain	4
S2576	ΔflaAB ₂	markerless in-frame deletion of the lateral flagellins (<i>sputcn32_3455-3456</i>)	5
S3297	ΔpdeB	markerless in-frame deletion of the gene <i>sputcn32_3405</i> (<i>pdeB</i>)	6
S4234	<i>pdeB-sfgfp</i>	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i>	6
S4237	<i>pdeB-E637A-sfgfp</i>	functional markerless substitution of the EIL motif to AIL (residue 637) in the background of <i>pdeB-sfgfp</i>	6
S4357	ΔflaAB ₂ ΔpdeB	markerless in-frame deletion of the gene <i>pdeB</i> (<i>sputcn32_3405</i>) in a background with deleted lateral flagellins	6
S6452	<i>pdeB-gfp</i> V522G V523G Q524G	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of Q524G	this study
S6453	<i>pdeB-gfp</i> K490G V491G M492G Q593G	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of M492G Q593G	this study
S6454	<i>pdeB-GFP</i> R557G A558G P559G Y560G	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of A558G P559G Y560G	this study
S6496	ΔflaAB ₂ ΔmshE	markerless in-frame deletion of the gene <i>mshE</i> (<i>sputcn32_0563</i>) in a background with deleted lateral flagellins	this study
S6497	ΔflaAB ₂ ΔpdeB ΔmshE	markerless in-frame deletion of the gene <i>mshE</i> (<i>sputcn32_0563</i>) in a background with deleted lateral flagellins and deletion of the gene <i>pdeB</i> (<i>sputcn32_3405</i>)	this study
S6527	<i>pdeB-gfp</i> K490D Q493A	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of K490D Q493A	this study
S6528	<i>pdeB-gfp</i> Q524A K527D Q528A	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of Q524A K527D Q528A	this study
S6659	<i>pdeB-gfp</i> Q524S Q528S	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of Q524S Q528S	this study
S6683	ΔflaAB ₂ mshA S68C	functional substitution of <i>mshA</i> S68C in a background with deleted lateral flagellins	this study
S6684	ΔflaAB ₂ ΔpdeB mshA S68C	functional substitution of <i>mshA</i> S68C in a background with deleted lateral flagellins and deletion of the gene <i>pdeB</i> (<i>sputcn32_3405</i>)	this study

S6688	$\Delta flaAB_2 \Delta aggA$	markerless in-frame deletion of the gene <i>aggA</i> (<i>sputcn32_3594</i>) in a background with deleted lateral flagellins	this study
S6689	$\Delta flaAB_2 \Delta pdeB \Delta aggA$	markerless in-frame deletion of the gene <i>aggA</i> (<i>sputcn32_3594</i>) in a background with deleted lateral flagellins and deletion of the gene <i>pdeB</i> (<i>sputcn32_3405</i>)	this study
S6690	$\Delta flaAB_2 \Delta pdeB \Delta mshE \Delta aggA$	markerless in-frame deletion of the gene <i>aggA</i> (<i>sputcn32_3594</i>) in a background with deleted lateral flagellins, and deletion of the genes <i>pdeB</i> (<i>sputcn32_3405</i>) and <i>mshE</i> (<i>sputcn32_0563</i>)	this study
S6691	$\Delta flaAB_2 \Delta pilB$	markerless in-frame deletion of the gene <i>pilB</i> (<i>sputcn32_3423</i>) in a background with deleted lateral flagellins	this study
S6692	$\Delta flaAB_2 \Delta pdeB \Delta pilB$	markerless in-frame deletion of the gene <i>pilB</i> (<i>sputcn32_3423</i>) in a background with deleted lateral flagellins and deletion of the gene <i>pdeB</i> (<i>sputcn32_3405</i>)	this study
S6693	$\Delta flaAB_2 \Delta pdeB \Delta mshE \Delta pilB$	markerless in-frame deletion of the gene <i>pilB</i> (<i>sputcn32_3423</i>) in a background with deleted lateral flagellins, and deletion of the genes <i>pdeB</i> (<i>sputcn32_3405</i>) and <i>mshE</i> (<i>sputcn32_0563</i>)	this study
S6729	<i>pdeB-gfp</i> K527E Q528S	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of K527E Q528S	this study
S7024	<i>pdeB-mvenus</i>	markerless in-frame fusion of <i>pdeB</i> with <i>mvenus</i>	this study
S7025	<i>pdeB-mvenus</i> D508A E509A	markerless in-frame fusion of <i>pdeB</i> D508A E509A with <i>mvenus</i>	this study
S7026	<i>pdeB-mvenus</i> E637A	markerless in-frame fusion of <i>pdeB</i> E637A with <i>mvenus</i>	this study
S7243	<i>pdeB-gfp</i> K527S Q528S	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of K527S Q528S	this study
S7244	<i>pdeB-gfp</i> K527S	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of K527S	this study
S7245	<i>pdeB-gfp</i> G497A	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of G497A	this study
S7246	<i>pdeB-gfp</i> Q499S	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of Q499S	this study
S7247	<i>pdeB-gfp</i> E500S	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of E500S	this study
S7444	$\Delta flaAB_2 \Delta mshE \Delta aggA$	markerless in-frame deletion of the gene <i>aggA</i> (<i>sputcn32_3594</i>) in a background with deleted lateral flagellins and deletion of the gene <i>mshE</i> (<i>sputcn32_0563</i>)	this study
S7445	$\Delta flaAB_2 \Delta mshE \Delta pilB$	markerless in-frame deletion of the gene <i>pilB</i> (<i>sputcn32_3423</i>) in a background with deleted lateral flagellins and deletion of the gene <i>mshE</i> (<i>sputcn32_0563</i>)	this study
S7504	<i>pdeB</i> K527E Q528S	markerless in-frame substitution of <i>pdeB</i> K527E Q528S	this study
S7505	<i>pdeB</i> G497A	markerless in-frame substitution of <i>pdeB</i> G497A	this study
S7506	<i>pdeB</i> K578S	markerless in-frame substitution of <i>pdeB</i> K578S	this study

S7507	<i>pdeB-mvenus</i> K527E Q528S	markerless in-frame fusion of <i>pdeB</i> -K527E Q528S with <i>mvenus</i>	this study
S7508	<i>pdeB</i> K580S	markerless in-frame substitution of <i>pdeB</i> K580S	this study
S7562	<i>pdeB-gfp</i> K578S	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of K578S	this study
S7564	<i>pdeB-gfp</i> K580S	markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> substitution of K580S	this study
S7614	pMMB-Gm-Bc3-5 AAV (hok-sok)	wildtype strain containing the c-di-GMP biosensor plasmid	⁷
S7616	$\Delta pdeB$ pMMB-Gm-Bc3-5 AAV (hok-sok)	c-di-GMP biosensor plasmid in the background of deleted <i>pdeB</i> (<i>sputcn32_3405</i>)	⁷
S7653	<i>pdeB</i> K527E Q528S pMMB-Gm-Bc3-5 AAV (hok-sok)	c-di-GMP biosensor plasmid in the background of <i>pdeB</i> K527E Q528S substitution	⁷
S7654	<i>pdeB</i> G497A pMMB-Gm-Bc3-5 AAV (hok-sok)	c-di-GMP biosensor plasmid in the background of <i>pdeB</i> G497A substitution	⁷
S7655	<i>pdeB</i> K578S pMMB-Gm-Bc3-5 AAV (hok-sok)	c-di-GMP biosensor plasmid in the background of <i>pdeB</i> K578S substitution	⁷
S7691	<i>lapA</i> -GS-3xFLAG	functional markerless in-frame tag of 3xFLAG to the C-terminus of <i>lapA</i> via a flexible GS-linker	this study
S7692	$\Delta pdeB$ <i>lapA</i> -GS-3xFLAG	functional markerless in-frame tag of 3xFLAG to the C-terminus of <i>lapA</i> via a flexible GS-linker in the background of deleted <i>pdeB</i> (<i>sputcn32_3405</i>)	this study
S7703	<i>lapB</i> -GS-3xFLAG	functional markerless in-frame tag of 3xFLAG to the C-terminus of <i>lapB</i> via a flexible GS-linker	this study
S7704	$\Delta pdeB$ <i>lapB</i> -GS-3xFLAG	functional markerless in-frame tag of 3xFLAG to the C-terminus of <i>lapB</i> via a flexible GS-linker in the background of deleted <i>pdeB</i> (<i>sputcn32_3405</i>)	this study

***Shewanella oneidensis* MR-1 strains**

S79	MR-1 wt	wildtype strain	⁸
S7296	$\Delta pdeB$	Markerless in-frame deletion of <i>pdeB</i> of <i>S. oneidensis</i> MR-1	this study
S7294	<i>pdeB-gfp</i>	Markerless in-frame fusion of <i>pdeB</i> with <i>sfgfp</i> in <i>S. oneidensis</i> MR-1	this study
S7423	pMMB-Gm-Bc3-5 AAV (hok-sok)	MR-1 wildtype strain containing the c-di-GMP biosensor plasmid	⁷
S7425	$\Delta pdeB$ pMMB-HS-Bc-3-5-AAV (hok-sok)	c-di-GMP biosensor plasmid in the background of deleted <i>pdeB</i> (SO_0437)	⁷

Supplementary Table 2: Plasmids that were used in this study

Plasmid	Relevant genotype or phenotype	Reference
pNPTS-138-R6KT	<i>mobRP4+</i> <i>ori-R6K</i> <i>sacB</i> β-galactosidase fragment alpha, suicide plasmid for in frame deletions/insertions in <i>Shewanella</i> , Km ^r	9
pET-24c	overproduction vector for His-tagged proteins	10
pBTOK	pBBR1-MCS2 backbone (pBBR origin, Km ^r); TetR, Promoter and multiple cloning site of pASK-IBA3plus and <i>E. coli</i> <i>rrnB1</i> T1 and lambda phage T0 terminator.	11
pMMB-Gm-Bc3-5 AAV (hok-sok)	Overproduction plasmid, inducible with anhydrotetracycline pMMB67EH (Gm) backbone containing the c-di-GMP biosensor (turboRFP with an AAV tag) and also the hok/sok region from pXB300. Used as c-di-GMP reporter.	7
overexpression vectors		
pET24c MBP-PdeB (MR-1) GGDEF-6xHis	Vector used to express the GGDEF-domain of MR-1 PdeB (residues 417 - 585) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c MBP-PdeB (MR-1) GGDEF-6xHis K524S	Vector used to express the GGDEF-domain of MR-1 PdeB (residues 417 - 585) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c MBP-PdeB (MR-1) GGDEF-6xHis Q525S	Vector used to express the GGDEF-domain of MR-1 PdeB (residues 417 - 585) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c MBP-PdeB (MR-1) GGDEF-6xHis K524E Q525S	Vector used to express the GGDEF-domain of MR-1 PdeB (residues 417 - 585) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c MBP-PdeB (MR-1) GGDEF-6xHis G494A	Vector used to express the GGDEF-domain of MR-1 PdeB (residues 417 - 585) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c MBP-PdeB (MR-1) GGDEF-6xHis E497S	Vector used to express the GGDEF-domain of MR-1 PdeB (residues 417 - 585) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c MBP-PdeB (MR-1) PAS-GGDEF-6xHis	Vector used to express the PAS- and GGDEF-domain of MR-1 PdeB (residues 304 - 585) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c MBP-PdeB (MR-1) PAS-GGDEF-6xHis K524E Q525S	Vector used to express the PAS- and GGDEF-domain of MR-1 PdeB (residues 304 - 585) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c (MR-1) FimV-Cdomain-6xHis	Vector used to express the C-terminal domain of MR-1 FimV (residues 1000 - 1110) with C-terminal 6xHis translational fusion	this study
pET24c 3xFLAG-(CN-32) HubP-FimV-Cdomain-6xHis	Vector used to express the C-terminal domain of CN-32 FimV (residues 1000 - 1110) with C-terminal 6xHis translational fusion and 3xFLAG	this study
pET24c MBP-PdeB (CN-32) GGDEF-EAL-6xHis	Vector used to express the GGDEF- and EAL-domain of CN-32 PdeB (residues 420-847) with N-terminal MBP and C-terminal 6xHis translational fusion	this study
pET24c <i>mshE</i> -6xHis	Vector used to express MshE of CN-32 with C-terminal 6xHis translational fusion	this study
pET24c <i>mshE</i> _Ndomain-6xHis	Vector used to express the N-terminal domain of CN-32 MshE (residues 2 - 145) with C-terminal 6xHis translational fusion	this study
pET24c <i>pilB</i> _Ndomain-6xHis	Vector used to express the N-terminal domain of CN-32 PilB (residues 2 - 145) with C-terminal 6xHis translational fusion	this study
pBTOK <i>dgcA</i> -6xHis	Vector for ectopical expression of <i>dgcA</i> (<i>E. coli</i>) in <i>S. putrefaciens</i> CN-32 with C-terminal 6xHis	this study
pBTOK <i>dgcA</i> -6xHis D216E	Vector for ectopical expression of <i>dgcA</i> (<i>E. coli</i>) in <i>S. putrefaciens</i> CN-32 with C-terminal 6xHis and D216E	this study

pBTOK <i>dgcA</i> -6xHis E276K	Vector for ectopical expression of <i>dgcA</i> (<i>E. coli</i>) in <i>S. putrefaciens</i> CN-32 with C-terminal 6xHis and E276K	this study
In-frame insertion vectors		
pNPTS CN-32 <i>pdeB-gfp</i> K527S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K527S	this study
pNPTS CN-32 <i>pdeB-gfp</i> K527S Q528S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K527S Q528S	this study
pNPTS CN-32 <i>pdeB-gfp</i> K527D	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K527D	this study
pNPTS CN-32 <i>pdeB-gfp</i> K527D Q528S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K527D Q528S	this study
pNPTS CN-32 <i>pdeB-gfp</i> Q524A K527D Q528A	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with Q524A K527D Q528A	this study
pNPTS CN-32 <i>pdeB-gfp</i> G497A	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with G497A	this study
pNPTS CN-32 <i>pdeB-gfp</i> Q499S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with Q499S	this study
pNPTS CN-32 <i>pdeB-gfp</i> E500S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with E500S	this study
pNPTS CN-32 <i>pdeB-gfp</i> Q528S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with Q528S	this study
pNPTS CN-32 <i>pdeB-gfp</i> Q524S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with Q524S	this study
pNPTS CN-32 <i>pdeB-gfp</i> Q524S Q528S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with Q524S Q528S	this study
pNPTS CN-32 <i>pdeB-gfp</i> K527E Q528S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K527E Q528S	this study
pNPTS CN-32 <i>pdeB-gfp</i> K490D Q493A	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K490D Q493A	this study
pNPTS CN-32 <i>pdeB-gfp</i> R557G A558G P559G Y560G	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with R557G A558G P559G Y560G	this study
pNPTS CN-32 <i>pdeB-gfp</i> V522G V523G Q524G	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with V522G V523G Q524G	this study
pNPTS CN-32 <i>pdeB-gfp</i> K490G V491G M492G Q593G	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K490G V491G M492G Q593G	this study
pNPTS CN-32 <i>pdeB-gfp</i> K578S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K578S	this study
pNPTS CN-32 <i>pdeB-gfp</i> K580S	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. putrefaciens</i> CN-32 with K580S	this study
pNPTS CN-32 <i>pdeB</i> G497A	Suicide vector for markerless in-frame insertion of <i>pdeB</i> of <i>S. putrefaciens</i> CN-32 with G497A	this study
pNPTS CN-32 <i>pdeB</i> K527E Q528S	Suicide vector for markerless in-frame insertion of <i>pdeB</i> of <i>S. putrefaciens</i> CN-32 with K527E Q528S	this study
pNPTS CN-32 <i>pdeB</i> K578S	Suicide vector for markerless in-frame insertion of <i>pdeB</i> of <i>S. putrefaciens</i> CN-32 with K578S	this study
pNPTS CN-32 <i>pdeB</i> K580S	Suicide vector for markerless in-frame insertion of <i>pdeB</i> of <i>S. putrefaciens</i> CN-32 with K580S	this study
pNPTS CN-32 <i>pdeB-venus</i>	Suicide vector for markerless in-frame insertion of <i>pdeB-venus</i> of <i>S. putrefaciens</i> CN-32	this study
pNPTS CN-32 <i>pdeB-venus</i> D508A E509A	Suicide vector for markerless in-frame insertion of <i>pdeB-venus</i> of <i>S. putrefaciens</i> CN-32 with 508A E509A	this study
pNPTS CN-32 <i>pdeB-venus</i> E637A	Suicide vector for markerless in-frame insertion of <i>pdeB-venus</i> of <i>S. putrefaciens</i> CN-32 with E637A	this study
pNPTS CN-32 <i>lapA-GS-3xFLAG</i>	Suicide vector for markerless in-frame insertion of 3xFLAG to the C-terminus of <i>lapA</i> via a flexible GS-linker	this study
pNPTS CN-32 <i>lapB-GS-3xFLAG</i>	Suicide vector for markerless in-frame insertion of 3xFLAG to the C-terminus of <i>lapB</i> via a flexible GS-linker	this study
pNPTS CN-32 <i>mshA</i> S68C	Suicide vector for markerless in-frame insertion of <i>mshA</i> of <i>S. putrefaciens</i> CN-32 with S68C	this study

pNPTS MR-1 <i>pdeB-gfp</i>	Suicide vector for markerless in-frame insertion of <i>pdeB-gfp</i> of <i>S. oneidensis</i> MR-1	this study
In-frame deletion vectors		
pNPTS CN-32 $\Delta mshE$	Suicide vector for markerless in-frame deletion of <i>mshE</i> of <i>S. putrefaciens</i> CN-32	this study
pNPTS CN-32 $\Delta aggA$	Suicide vector for markerless in-frame deletion of <i>aggA</i> of <i>S. putrefaciens</i> CN-32	this study
pNPTS CN-32 $\Delta pilB$	Suicide vector for markerless in-frame deletion of <i>pilB</i> of <i>S. putrefaciens</i> CN-32	this study
pNPTS MR-1 $\Delta pdeB$	Suicide vector for markerless in-frame deletion of <i>pdeB</i> of <i>S. oneidensis</i> MR-1	this study

Supplementary Table 3: Primer that were used in this study

Plasmid	Primer	Sequence
pET24c MBP-PdeB (MR-1) GGDEF-6xHis	TR258 MBP fw	TTAACCTTAAGAAGGAGATATAATGA AAATAGAAGAAGGTAAACTGGTAATCTG G
	TR259 MBP rv	GCTCCCCCGAGGTTGTTATTGTTA TTGT
	TR260	AATAACAACACCTCGGGGGCAGCGAA GAACCTTCTTAAGCATCAGCTAC
	TR257	GTGGTGGTGGTGGTGGTCAATGGT GATGGTGATGGTGGTAAATGTGGATTG GTTGGTGC
pET24c MBP-PdeB (MR-1) GGDEF-6xHis K524S	TR510 MBP ol plas fw	TTAACCTTAAGAAGGAGATATAATGA AAATAGAAGAAGGTAAACTGGTAATCTG G
	TR511 So KtoS fw	CAATAATTGGCTCAGCACTGCGCCAC AGC
	TR512 So KtoS rv	GTTGCTGAGCCAATTATTGCTCAAGTA TCGCTGC
	TR513 soGGDEF ol plas rv	GTGGTGGTGGTGGTGGTCAATGGT GATGGTGATGGTGGTAAATGTGGATTG GTTGGTGC
pET24c MBP-PdeB (MR-1) GGDEF-6xHis Q525S	TR510 MBP ol plas fw	TTAACCTTAAGAAGGAGATATAATGA AAATAGAAGAAGGTAAACTGGTAATCTG G
	TR514 So QtoS fw	GAGCAATAATGCTCTTCAGCACTGCGC CACAGC
	TR515 So QtoS rv	GCTGAAGAGCATTATTGCTCAAGTATCG CTGC
	TR513 soGGDEF ol plas rv	GTGGTGGTGGTGGTGGTCAATGGT GATGGTGATGGTGGTAAATGTGGATTG GTTGGTGC
pET24c MBP-PdeB (MR-1) GGDEF-6xHis K524E Q525S	TR510 MBP ol plas fw	TTAACCTTAAGAAGGAGATATAATGA AAATAGAAGAAGGTAAACTGGTAATCTG G
	TR593 SO KQ to ES rv	TAATGCTTCCAGCACTGCGCCACAGC TAA
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	TR513 soGGDEF ol plas rv	GTGGTGGTGGTGGTGGTCAATGGT GATGGTGATGGTGGTAAATGTGGATTG GTTGGTGC

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	TR555 G494A rv	ATGTCTTGCGCCACAGGAATTATTAGCC CGCA
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	TR383	GTGGTGGTGGTGGTGGTGCCTAACGAC GATAAAGATTATCAAAGGCC
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	VK233 EcoRV 3591- down rv	GCCAAGCTTCTGCAGGATGGCTTCTA GTGACTCAATATTGAGTGTC
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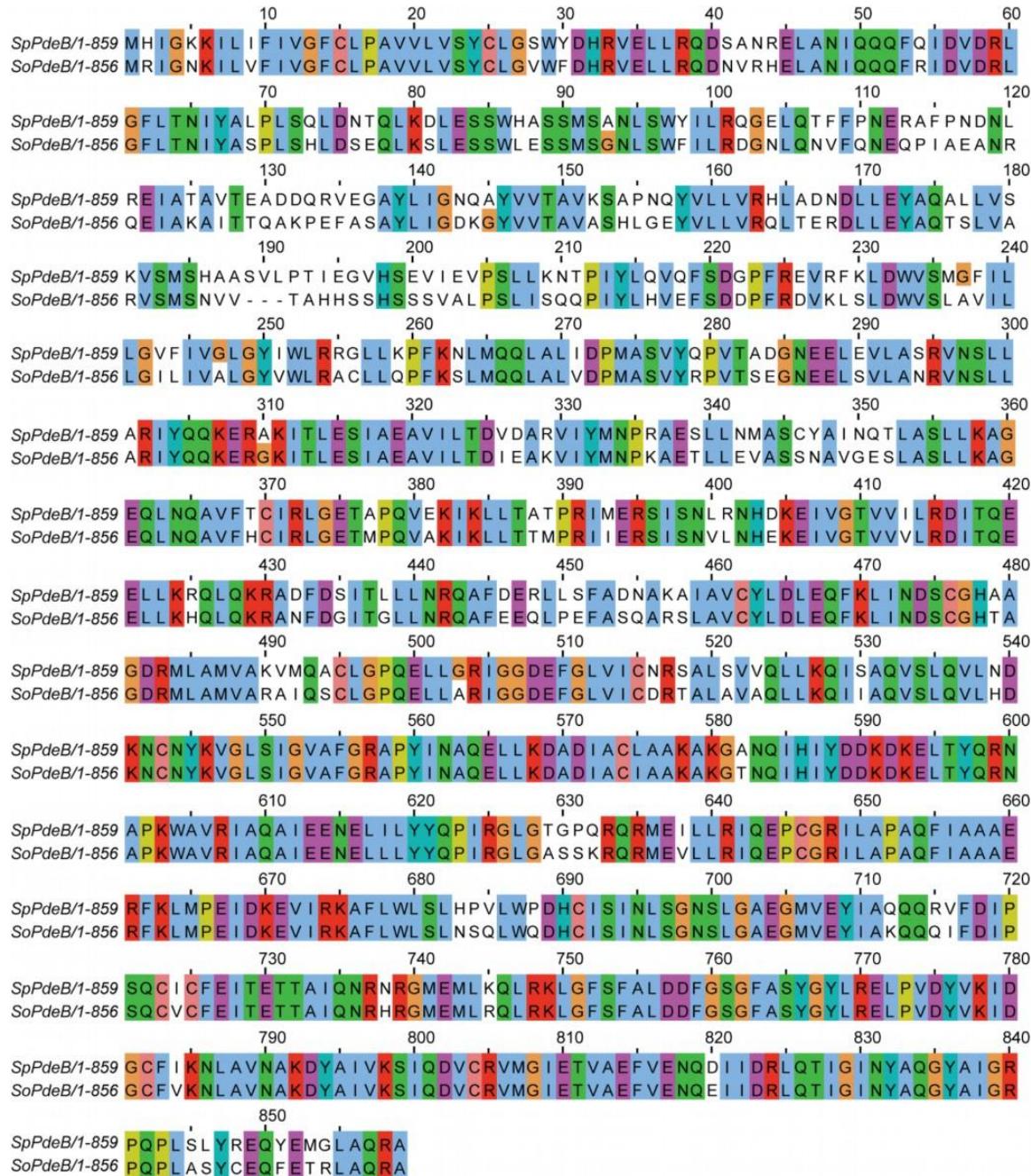
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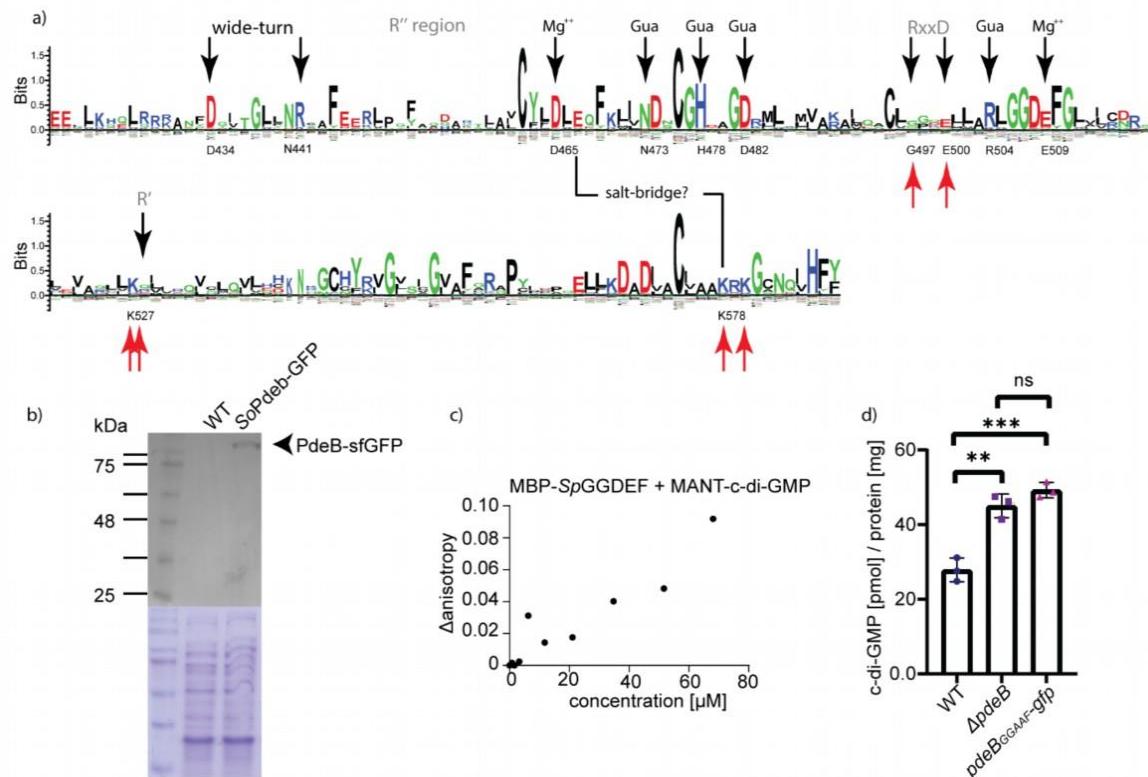
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Supplementary References

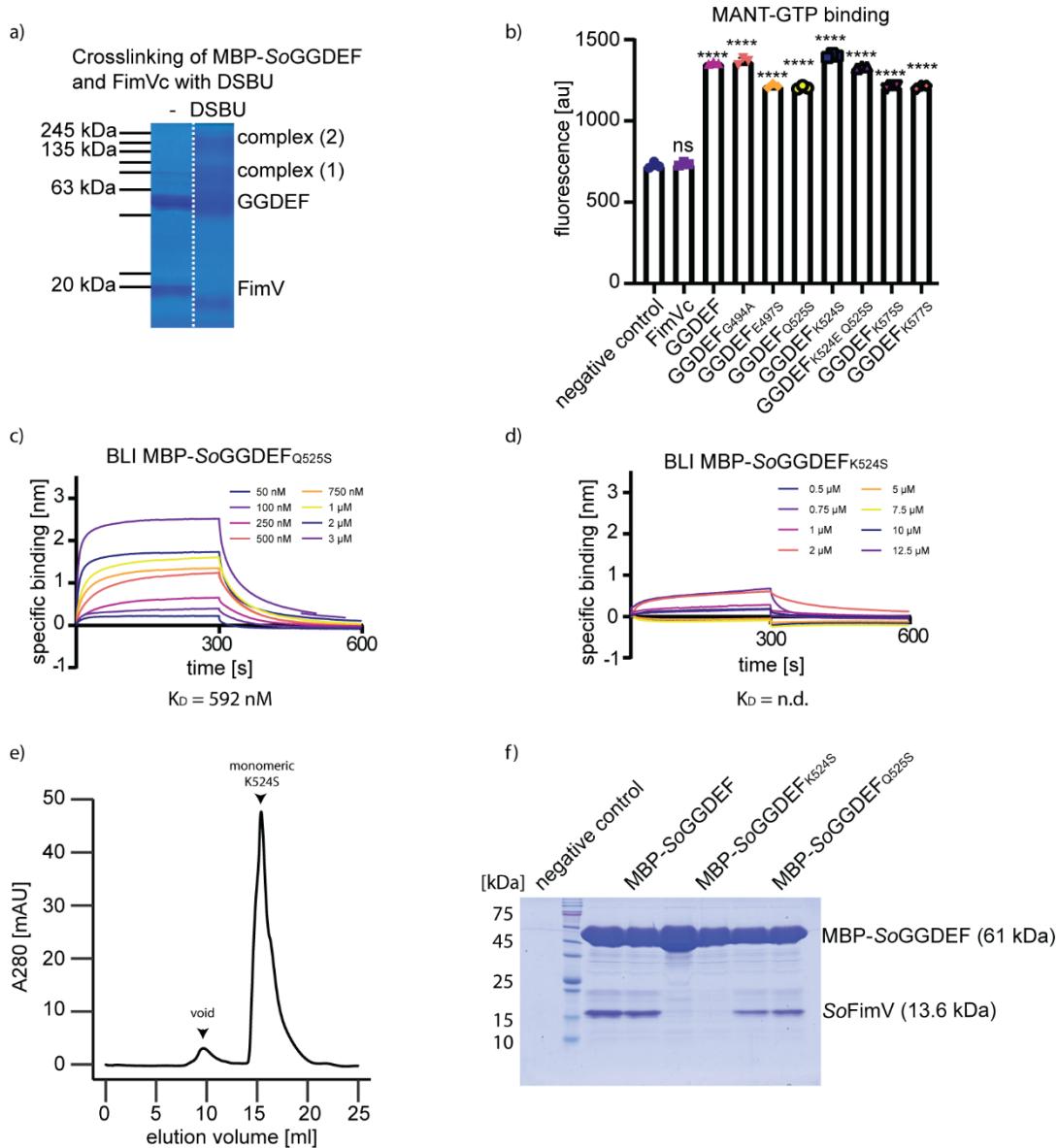
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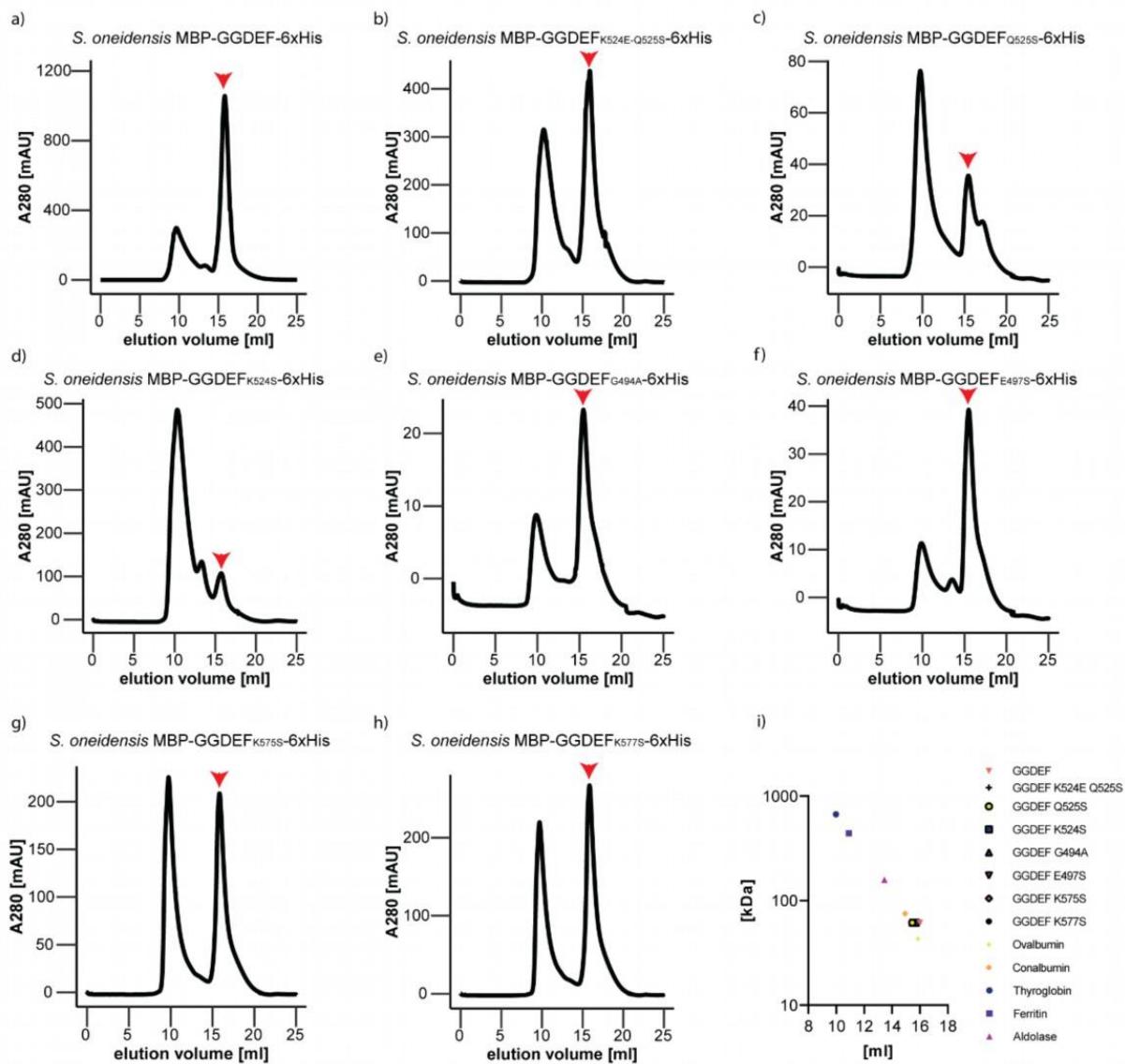
Supplementary Figure 1: Alignment of SpPdeB with SoPdeB. The conservation between PdeB homologous of *S. putrefaciens* and *S. oneidensis* was analyzed by aligning the amino acid sequences. The two proteins share 79% identity. Colored residues indicate 100% conservation between the two proteins. Periplasmic region: 30 - 230; HAMP: 255 - 308; PAS: 308 - 376; GGDEF: 420 - 588; EAL: 598 - 846.



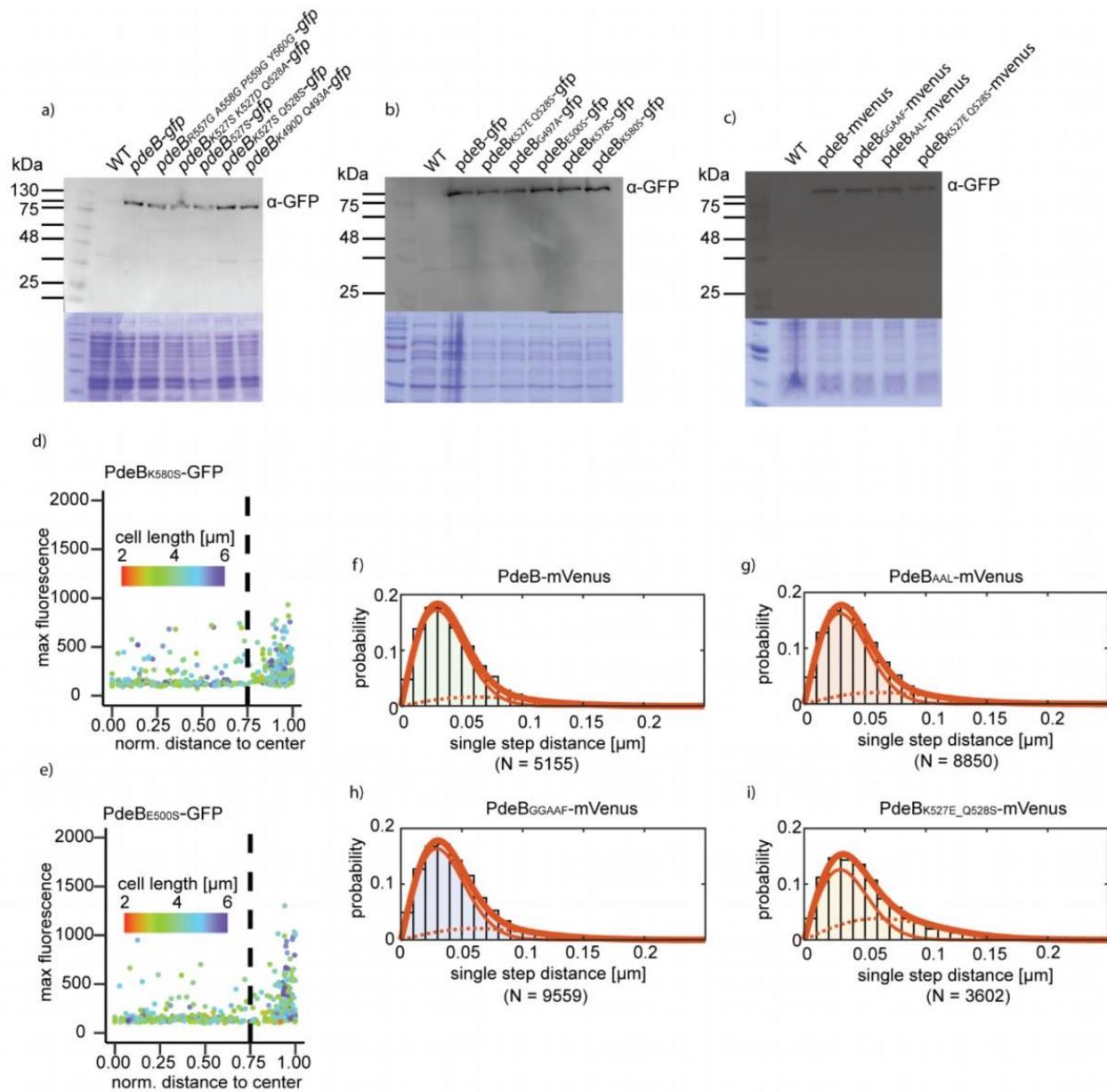
Supplementary Figure 2. a) Position-based weight map of the GGDEF domain of 50 *PdeB* homologues from different *Shewanella* species. Characteristic features of GGDEF domains are marked with black arrows, and degenerated or missing motifs are indicated in gray. Residues that are important for the GGDEF_{PdeB}-FimV_{HubP} interaction are highlighted by red arrows. **b)** The stability and expression of genomic SoPdeB-sfGFP fusions was verified by immunoblot analysis. **c)** The MANT-c-di-GMP binding of the GGDEF domain of *SoPdeB* was tested by fluorescence anisotropy assays. No binding curve was observed, but only unspecific binding at unphysiological high ligand concentrations. **d)** The effect of GTP binding to GGDEF_{PdeB} on the PDE activity of PdeB was determined by introducing mutations the GGDEF motif. The cellular c-di-GMP content was then extracted and quantified by MS. The single point mutation within the GGDEF motif results in a similar increase in cellular c-di-GMP as deletion of *pdeB*. Significance was tested by using the unpaired t-test (* P < 0.05, ** P < 0.005, *** P < 0.0005).



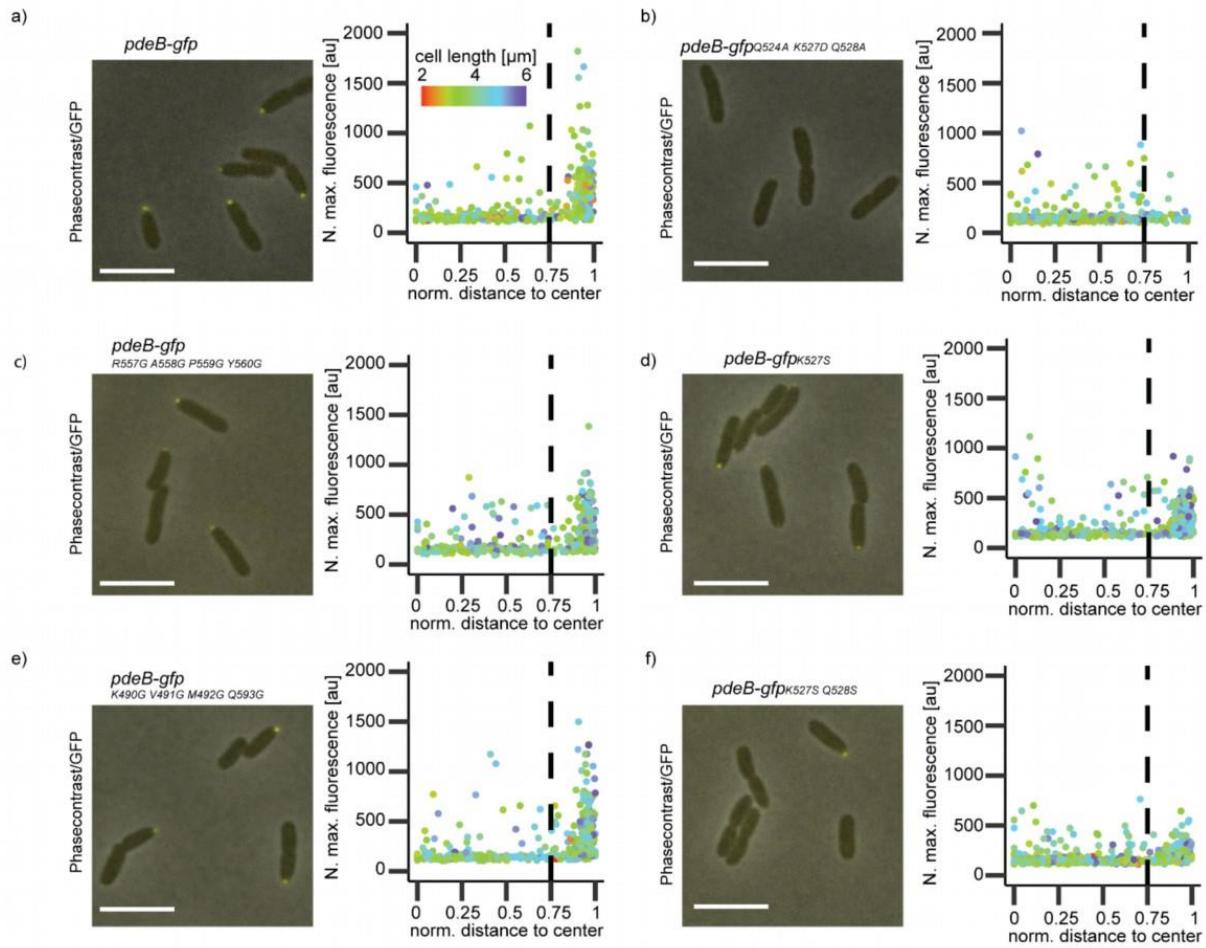
Supplementary Figure 3: Structural and sequential features of PdeB_{GGDEF} and HubP_{FimV}. **a)** The crosslinking of SoGGDEF with SoHubP was verified by SDS-PAGE. The unprocessed PAGE can be found in Supplementary Figure 11. **b)** Functionality of MBP-SoGGDEF proteins used for BLI was shown by MANT-GTP binding assays. All mutated versions are able to bind MANT-GTP, as indicated by the increased fluorescence. Significance was tested by using the unpaired t-test (* P < 0.05, ** P < 0.005, *** P < 0.0005, **** P < 0.00005). **c, d)** BLI assays for GGDEF proteins with substitutions in the R' I-site show decreased affinity to FimV compared to the wild type. The purified MBP-GGDEF_{K524S} showed aggregation upon production and unspecific binding in BLI assays and was therefore not suitable for determination of exact KD values. **e)** The stability was therefore tested by storing the protein for three days at 4°C and further analysed with SEC. The sample remained mostly in monomeric form. **f)** The interaction of MBP-GGDEF_{K524S} with FimV was tested by pull-down assays where the wild-type version and MBP-GGDEF_{Q525S} served as controls. No binding was observed when K524 was mutated.



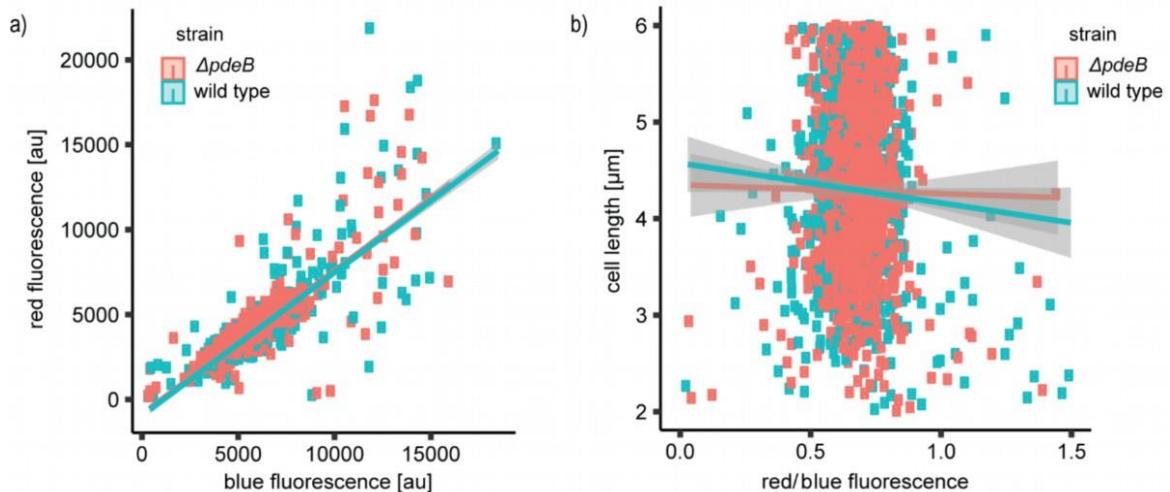
Supplementary Figure 4: SEC of GGDEF domains. **a-h)** The mutated GGDEF_{PdeB} domains of *S. oneidensis* were purified as MBP-fusion proteins. The chromatograms are shown in a-h. The elution volume in ml is plotted against the absorbance at 280 nm. Red arrows indicate the peak for the monomeric proteins of interest. **i)** The elution volume of the proteins of interests was plotted against the molecular weight in kDa, together with globular proteins included in the high molecular weight calibration kit (GE healthcare). All GGDEF proteins elute at roughly the same elution volume, indicating that the introduced amino acid substitutions do not alter the mutant proteins' structure.



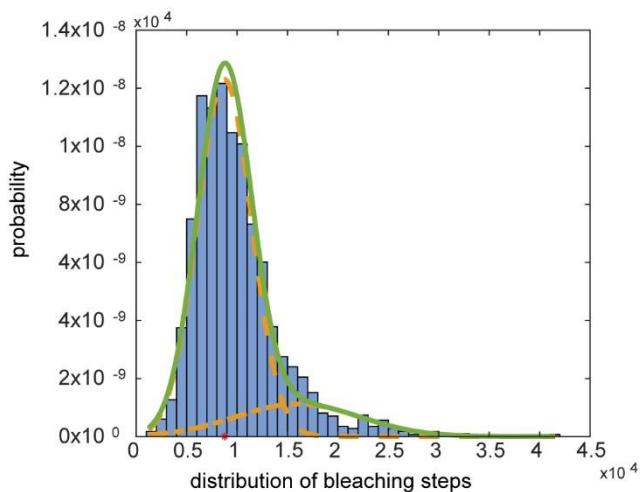
Supplementary Figure 5. a-b) Stability and expression of *SpPdeB*-sfGFP mutants were verified using Western blot analysis. **c)** Stability of *SpPdeB*-mVenus used for single molecule microscopy was also shown by Western blot analysis. **d-e)** Scatterplots of fluorescence microscopy of mutants with reduced PdeB-sfGFP localization, displaying the normalized distance from the central plane of the cell. Both mutations reduce the polar localization of PdeB-sfGFP, likely due to reduced affinity to HubP. **f-i)** Fits for the jump distance analyses used for the data of the bubble blot. Solid thin line Rayleigh fit for the slow population, dotted line fast population, and thick solid line combination of both fits. R^2 of values all fits were higher than 0.999. For all mVenus fusions, a two population-fit was the best.



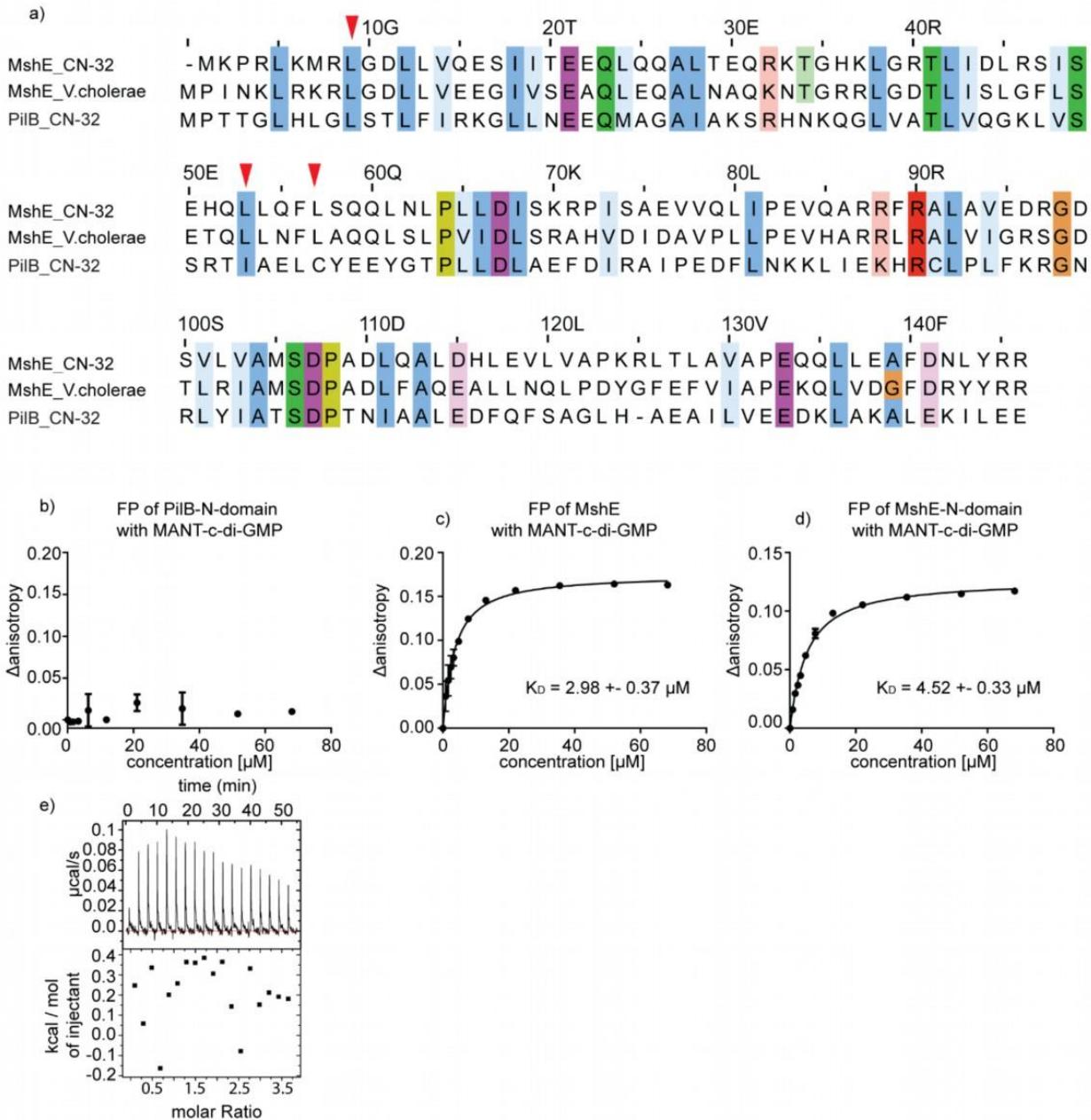
Supplementary Figure 6: Screening for residues involved in the polar localization of PdeB. **a-f)** Residues at different regions in the GGDEF domain of *SpPdeB-sfGFP* were genomically mutated and localization behavior was observed using fluorescence microscopy. The localization behaviors of mutants are shown as scatter plots displaying the normalized distance from the central plane of the cell, wild-type *SpPdeB-sfGFP* serves as control. Mutating residues of the R' site (b, d, f) leads to reduced polar localization, while substitutions in the other two regions only had minor effects.



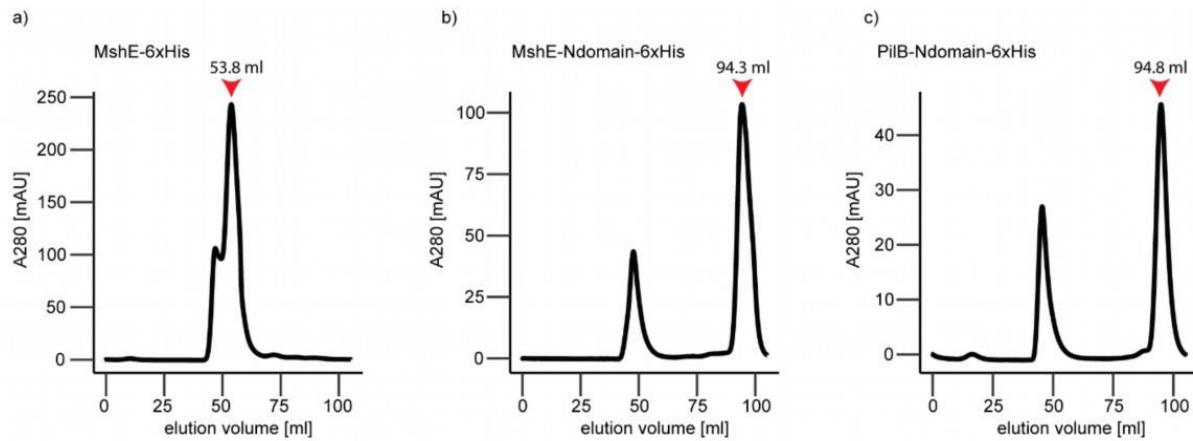
Supplementary Figure 7: c-di-GMP single cell reporter controls. **a)** The functionality of the fluorescence based c-di-GMP reporter was tested for *S. putrefaciens* CN-32 by plotting the blue against the red fluorescence and testing for linear correlation in presence and absence of *pdeB*. **b)** A correlation of cell length with the c-di-GMP level was tested by plotting the quotient of the red fluorescence divided by the blue fluorescence against the cell length. No such correlation was found.



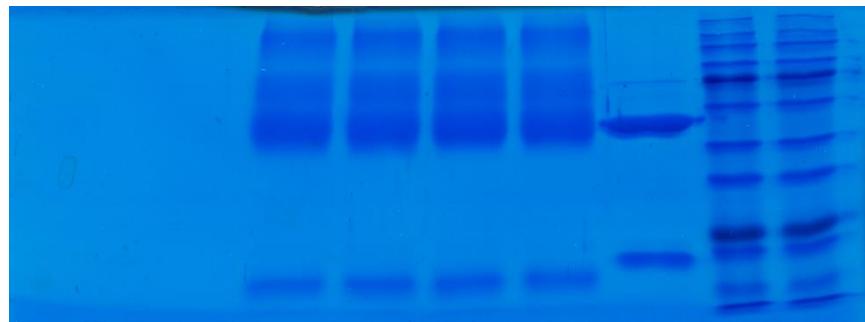
Supplementary Figure 8: Fluorescence-based molecule quantification of PdeB-mVenus. The distribution of bleaching steps within the movies is shown as histogram.



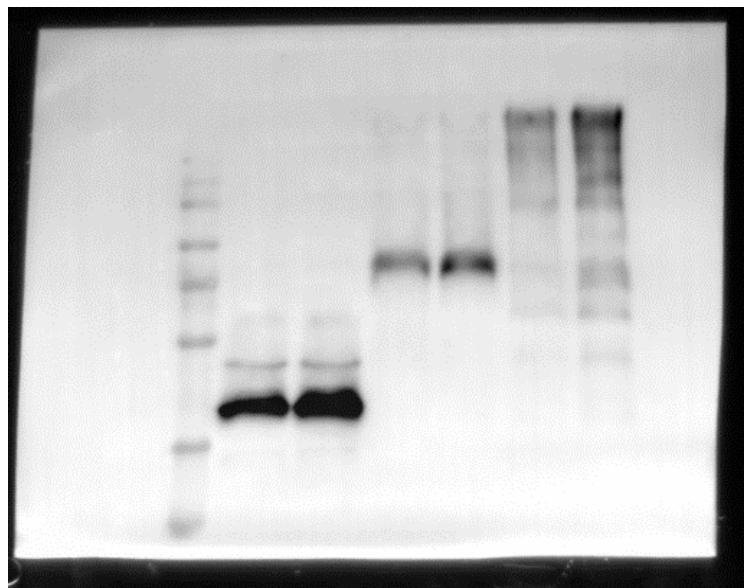
Supplementary Figure 9. **a)** Alignment of *PaMshE* with *SpMshE* and *SpPilB*. Leucines that are involved in c-di-GMP binding are marked with red arrows. **b-d)** The MANT-c-di-GMP binding of the N-terminal domain of PilB (b), MshE (c) and the N-terminal domain of MshE (d) was tested by fluorescence anisotropy measurements. Both MshE versions are able to bind MANT-c-di-GMP with high affinity, while PilB does not. **e)** The non-binding of c-di-GMP to PilB was verified by ITC.



Supplementary Figure 10: SEC of MshE and PilB. a-b) Chromatograms of the size-exclusion chromatography of the extension ATPase MshE. Chromatograms show the elution volume against the absorbance at 280 nm. Peaks that contain the protein of interest are indicated by red arrows. The full length MshE protein (a) eluted at 53.8 ml, indicating an oligomeric state (penta- or hexameric), while the N-terminal domain eluted as monomer. **c)** The N-terminal domain of PilB eluted as monomer.



Supplementary Figure 11: Uncropped scan of the crosslinking control PAGE shown in **Supplementary Figure 3a**. The first four panels from the left show different crosslinking preparations, followed by a control of non-crosslinked protein and two protein ladder standards to the right.



Supplementary Figure 12: Uncropped Western Blot shown in **Figure 6f**. The proteins in lane 2 and 3 right next to the protein standard belong to a different protein and were therefore removed.